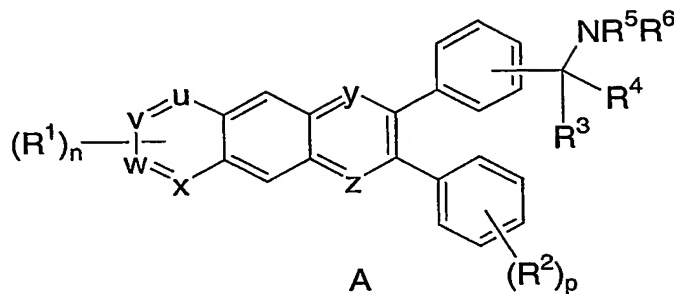


WHAT IS CLAIMED IS:

1. A compound of the Formula A:



5 wherein:

a is 0 or 1;

b is 0 or 1;

m is 0, 1 or 2;

10 n is 0, 1 or 2;

p is 0, 1, 2 or 3;

r is 0 or 1;

s is 0 or 1;

t is 2, 3, 4, 5 or 6;

15

u, v and x are independently selected from: CH and N;

w is selected from a bond, CH and N;

20 y and z are independently selected from: CH and N, provided that at least one of y and z is N;

R¹ is independently selected from:

- 25
- 1) (C=O)_aO_bC₁-C₁₀ alkyl,
 - 2) (C=O)_aO_baryl,
 - 3) C₂-C₁₀ alkenyl,
 - 4) C₂-C₁₀ alkynyl,
 - 5) (C=O)_aO_b heterocyclyl,

- 5 6) $(\text{C}=\text{O})_a\text{O}_b\text{C}_3\text{-C}_8$ cycloalkyl,
 7) CO_2H ,
 8) halo,
 9) CN,
 10) OH,
 11) $\text{O}_b\text{C}_1\text{-C}_6$ perfluoroalkyl,
 12) $\text{O}_a(\text{C}=\text{O})_b\text{NR}^7\text{R}^8$,
 13) $\text{NR}^c(\text{C}=\text{O})\text{NR}^7\text{R}^8$,
 14) $\text{S}(\text{O})_m\text{R}^a$,
 10 15) $\text{S}(\text{O})_2\text{NR}^7\text{R}^8$,
 16) $\text{NR}^c\text{S}(\text{O})_m\text{R}^a$,
 17) oxo,
 18) CHO,
 19) NO_2 ,
 15 20) $\text{NR}^c(\text{C}=\text{O})\text{O}_b\text{R}^a$,
 21) $\text{O}(\text{C}=\text{O})\text{O}_b\text{C}_1\text{-C}_{10}$ alkyl,
 22) $\text{O}(\text{C}=\text{O})\text{O}_b\text{C}_3\text{-C}_8$ cycloalkyl,
 23) $\text{O}(\text{C}=\text{O})\text{O}_b$ aryl, and
 24) $\text{O}(\text{C}=\text{O})\text{O}_b$ -heterocycle,
 20 said alkyl, aryl, alkenyl, alkynyl, heterocyclyl, and cycloalkyl optionally substituted
 with one or more substituents selected from R^Z ;

R^2 is independently selected from:

- 25 1) $(\text{C}=\text{O})_a\text{O}_b\text{C}_1\text{-C}_{10}$ alkyl,
 2) $(\text{C}=\text{O})_a\text{O}_b$ aryl,
 3) $\text{C}_2\text{-C}_{10}$ alkenyl,
 4) $\text{C}_2\text{-C}_{10}$ alkynyl,
 5) $(\text{C}=\text{O})_a\text{O}_b$ heterocyclyl,
 6) $(\text{C}=\text{O})_a\text{O}_b\text{C}_3\text{-C}_8$ cycloalkyl,
 30 7) CO_2H ,
 8) halo,
 9) CN,
 10) OH,
 11) $\text{O}_b\text{C}_1\text{-C}_6$ perfluoroalkyl,

- 12) $O_a(C=O)_bNR^7R^8$,
 13) $NR^c(C=O)NR^7R^8$,
 14) $S(O)_mR^a$,
 15) $S(O)_2NR^7R^8$,
 5 16) $NR^cS(O)_mR^a$,
 17) CHO ,
 18) NO_2 ,
 19) $NR^c(C=O)O_bR^a$,
 20) $O(C=O)O_bC_1-C_{10}$ alkyl,
 10 21) $O(C=O)O_bC_3-C_8$ cycloalkyl,
 22) $O(C=O)O_b$ aryl, and
 23) $O(C=O)O_b$ -heterocycle,

said alkyl, aryl, alkenyl, alkynyl, heterocyclyl, and cycloalkyl optionally substituted with one, two or three substituents selected from R^Z ;

15

R^3 and R^4 are independently selected from: H, C_1-C_6 -alkyl and C_1-C_6 -perfluoroalkyl, or

20

R^3 and R^4 are combined to form $-(CH_2)_t-$ wherein one of the carbon atoms is optionally replaced by a moiety selected from O, $S(O)_m$, $-N(R^b)C(O)-$, and $-N(COR^a)-$;

R^5 and R^6 are independently selected from:

- 25 1) H,
 2) $(C=O)O_bR^a$,
 3) C_1-C_{10} alkyl,
 4) aryl,
 5) C_2-C_{10} alkenyl,
 6) C_2-C_{10} alkynyl,
 30 7) heterocyclyl,
 8) C_3-C_8 cycloalkyl,
 9) SO_2R^a , and
 10) $(C=O)NR^b_2$,

said alkyl, cycloalkyl, aryl, heterocyclyl, alkenyl, and alkynyl is optionally substituted with one or more substituents selected from R^Z , or

5 R^5 and R^6 can be taken together with the nitrogen to which they are attached to form a monocyclic or bicyclic heterocycle with 5-7 members in each ring and optionally containing, in addition to the nitrogen, one or two additional heteroatoms selected from N, O and S, said monocyclic or bicyclic heterocycle optionally substituted with Q and also optionally substituted with one or more substituents selected from R^Z ;

10 R^7 and R^8 are independently selected from:

- 1) H,
- 2) $(C=O)O_bC_1-C_{10}$ alkyl,
- 3) $(C=O)O_bC_3-C_8$ cycloalkyl,
- 4) $(C=O)O_b$ aryl,
- 15 5) $(C=O)O_b$ heterocyclyl,
- 6) C_1-C_{10} alkyl,
- 7) aryl,
- 8) C_2-C_{10} alkenyl,
- 9) C_2-C_{10} alkynyl,
- 20 10) heterocyclyl,
- 11) C_3-C_8 cycloalkyl,
- 12) SO_2R^a , and
- 13) $(C=O)NR^b_2$,

25 said alkyl, cycloalkyl, aryl, heterocyclyl, alkenyl, and alkynyl is optionally substituted with one or more substituents selected from R^Z , or

R^7 and R^8 can be taken together with the nitrogen to which they are attached to form a monocyclic or bicyclic heterocycle with 5-7 members in each ring and optionally containing, in addition to the nitrogen, one or two additional heteroatoms selected from N, O and S, said monocyclic or bicyclic heterocycle optionally substituted with one or more substituents selected from R^Z ;

R^Z is selected from:

- 1) $(C=O)_rO_s(C_1-C_{10})$ alkyl,
- 35 2) $O_r(C_1-C_3)$ perfluoroalkyl,

- 3) (C₀-C₆)alkylene-S(O)_mR^a,
- 4) oxo,
- 5) OH,
- 6) halo,
- 5 7) CN,
- 8) (C=O)_rO_s(C₂-C₁₀)alkenyl,
- 9) (C=O)_rO_s(C₂-C₁₀)alkynyl,
- 10) (C=O)_rO_s(C₃-C₆)cycloalkyl,
- 11) (C=O)_rO_s(C₀-C₆)alkylene-aryl,
- 10 12) (C=O)_rO_s(C₀-C₆)alkylene-heterocyclyl,
- 13) (C=O)_rO_s(C₀-C₆)alkylene-N(R^b)₂,
- 14) C(O)R^a,
- 15) (C₀-C₆)alkylene-CO₂R^a,
- 16) C(O)H,
- 15 17) (C₀-C₆)alkylene-CO₂H,
- 18) C(O)N(R^b)₂,
- 19) S(O)_mR^a,
- 20) S(O)₂N(R^b)₂,
- 21) NR^c(C=O)O_bR^a,
- 20 22) O(C=O)O_bC₁-C₁₀ alkyl,
- 23) O(C=O)O_bC₃-C₈ cycloalkyl,
- 24) O(C=O)O_baryl, and
- 25) O(C=O)O_b-heterocycle,

25 said alkyl, alkenyl, alkynyl, cycloalkyl, aryl, and heterocyclyl is optionally substituted with up to three substituents selected from R^b, OH, (C₁-C₆)alkoxy, halogen, CO₂H, CN, O(C=O)C₁-C₆ alkyl, oxo, and N(R^b)₂;

30 R^a is substituted or unsubstituted (C₁-C₆)alkyl, substituted or unsubstituted (C₂-C₆)alkenyl, substituted or unsubstituted (C₂-C₆)alkynyl, substituted or unsubstituted (C₃-C₆)cycloalkyl, substituted or unsubstituted aryl, (C₁-C₆)perfluoroalkyl, 2,2,2-trifluoroethyl, or substituted or unsubstituted heterocyclyl; and

R^b is H, (C₁-C₆)alkyl, substituted or unsubstituted aryl, substituted or unsubstituted benzyl, substituted or unsubstituted heterocyclyl, (C₃-C₆)cycloalkyl, (C=O)OC₁-C₆ alkyl, (C=O)C₁-C₆ alkyl or S(O)₂R^a;

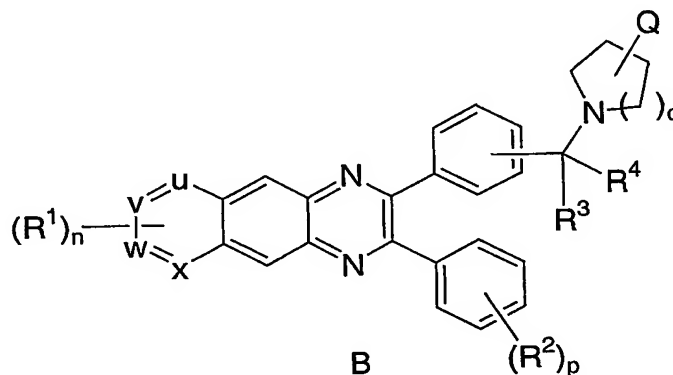
5 R^c is selected from:

- 1) H,
- 2) C₁-C₁₀ alkyl,
- 3) aryl,
- 4) C₂-C₁₀ alkenyl,
- 10 5) C₂-C₁₀ alkynyl,
- 6) heterocyclyl,
- 7) C₃-C₈ cycloalkyl,
- 8) C₁-C₆ perfluoroalkyl,

15 said alkyl, cycloalkyl, aryl, heterocyclyl, alkenyl, and alkynyl is optionally substituted with one or more substituents selected from R^z, or

or a pharmaceutically acceptable salt or a stereoisomer thereof.

2. A compound of the Formula B:



20

wherein:

- a is 0 or 1;
- b is 0 or 1;
- 25 m is 0, 1 or 2;
- n is 0, 1 or 2;

p is 0, 1 or 2;

q is 0, 1, 2, 3 or 4;

r is 0 or 1;

s is 0 or 1;

5 t is 2, 3, 4, 5 or 6;

u, v and x are independently selected from: CH and N;

w is selected from a bond, CH and N;

10

Q is selected from: -NR⁷R⁸, aryl and heterocyclyl, said aryl and heterocyclyl optionally substituted with one to three substituents selected from R^Z;

R¹ is independently selected from:

- | | |
|----|---|
| 15 | 1) (C=O) _a O _b C ₁ -C ₁₀ alkyl, |
| | 2) (C=O) _a O _b aryl, |
| | 3) C ₂ -C ₁₀ alkenyl, |
| | 4) C ₂ -C ₁₀ alkynyl, |
| | 5) (C=O) _a O _b heterocyclyl, |
| 20 | 6) (C=O) _a O _b C ₃ -C ₈ cycloalkyl, |
| | 7) CO ₂ H, |
| | 8) halo, |
| | 9) CN, |
| | 10) OH, |
| 25 | 11) O _b C ₁ -C ₆ perfluoroalkyl, |
| | 12) O _a (C=O) _b NR ⁷ R ⁸ , |
| | 13) NR ^c (C=O)NR ⁷ R ⁸ , |
| | 14) S(O) _m R ^a , |
| | 15) S(O) ₂ NR ⁷ R ⁸ , |
| 30 | 16) NR ^c S(O) _m R ^a , |
| | 17) oxo, |
| | 18) CHO, |
| | 19) NO ₂ , |
| | 20) NR ^c (C=O)O _b R ^a , |
| 35 | 21) O(C=O)O _b C ₁ -C ₁₀ alkyl, |

- 22) $O(C=O)O_bC_3-C_8$ cycloalkyl,
- 23) $O(C=O)O_b$ aryl, and
- 24) $O(C=O)O_b$ -heterocycle,

said alkyl, aryl, alkenyl, alkynyl, heterocyclyl, and cycloalkyl optionally substituted

5 with one or more substituents selected from R^Z ;

R^2 is independently selected from:

- 1) $(C=O)_aO_bC_1-C_{10}$ alkyl,
- 2) $(C=O)_aO_b$ aryl,
- 10 3) C_2-C_{10} alkenyl,
- 4) C_2-C_{10} alkynyl,
- 5) $(C=O)_aO_b$ heterocyclyl,
- 6) $(C=O)_aO_bC_3-C_8$ cycloalkyl,
- 7) CO_2H ,
- 15 8) halo,
- 9) CN ,
- 10) OH ,
- 11) $O_bC_1-C_6$ perfluoroalkyl,
- 12) $O_a(C=O)_bNR^7R^8$,
- 20 13) $NR^c(C=O)NR^7R^8$,
- 14) $S(O)_mR^a$,
- 15) $S(O)_2NR^7R^8$,
- 16) $NR^cS(O)_mR^a$,
- 17) CHO ,
- 25 18) NO_2 ,
- 19) $NR^c(C=O)O_bR^a$,
- 20) $O(C=O)O_bC_1-C_{10}$ alkyl,
- 21) $O(C=O)O_bC_3-C_8$ cycloalkyl,
- 22) $O(C=O)O_b$ aryl, and
- 30 23) $O(C=O)O_b$ -heterocycle,

said alkyl, aryl, alkenyl, alkynyl, heterocyclyl, and cycloalkyl optionally substituted with one, two or three substituents selected from R^Z ;

R^3 and R^4 are independently selected from: H, C_1-C_6 -alkyl and C_1-C_6 -perfluoroalkyl, or

R³ and R⁴ are combined to form -(CH₂)_t- wherein one of the carbon atoms is optionally replaced by a moiety selected from O, S(O)_m, -N(R^b)C(O)-, and -N(COR^a)-;

5

R⁷ and R⁸ are independently selected from:

- 1) H,
- 2) (C=O)O_bC₁-C₁₀ alkyl,
- 3) (C=O)O_bC₃-C₈ cycloalkyl,
- 10 4) (C=O)O_baryl,
- 5) (C=O)O_bheterocyclyl,
- 6) C₁-C₁₀ alkyl,
- 7) aryl,
- 8) C₂-C₁₀ alkenyl,
- 15 9) C₂-C₁₀ alkynyl,
- 10) heterocyclyl,
- 11) C₃-C₈ cycloalkyl,
- 12) SO₂R^a, and
- 13) (C=O)NR^b₂,

20 said alkyl, cycloalkyl, aryl, heterocyclyl, alkenyl, and alkynyl is optionally substituted with one or more substituents selected from R^z, or

R⁷ and R⁸ can be taken together with the nitrogen to which they are attached to form a monocyclic or bicyclic heterocycle with 5-7 members in each ring and optionally
25 containing, in addition to the nitrogen, one or two additional heteroatoms selected from N, O and S, said monocyclic or bicyclic heterocycle optionally substituted with one or more substituents selected from R^z;

R^z is selected from:

- 30 1) (C=O)_rO_s(C₁-C₁₀)alkyl,
- 2) O_r(C₁-C₃)perfluoroalkyl,
- 3) (C₀-C₆)alkylene-S(O)_mR^a,
- 4) oxo,
- 5) OH,
- 35 6) halo,

- 7) CN,
- 8) $(C=O)_rO_s(C_2-C_{10})$ alkenyl,
- 9) $(C=O)_rO_s(C_2-C_{10})$ alkynyl,
- 10) $(C=O)_rO_s(C_3-C_6)$ cycloalkyl,
- 5 11) $(C=O)_rO_s(C_0-C_6)$ alkylene-aryl,
- 12) $(C=O)_rO_s(C_0-C_6)$ alkylene-heterocyclyl,
- 13) $(C=O)_rO_s(C_0-C_6)$ alkylene- $N(R^b)_2$,
- 14) $C(O)R^a$,
- 15) (C_0-C_6) alkylene- CO_2R^a ,
- 10 16) $C(O)H$,
- 17) (C_0-C_6) alkylene- CO_2H ,
- 18) $C(O)N(R^b)_2$,
- 19) $S(O)_mR^a$,
- 20) $S(O)_2N(R^b)_2$,
- 15 21) $NR^c(C=O)O_bR^a$,
- 22) $O(C=O)O_bC_1-C_{10}$ alkyl,
- 23) $O(C=O)O_bC_3-C_8$ cycloalkyl,
- 24) $O(C=O)O_b$ aryl, and
- 25) $O(C=O)O_b$ -heterocycle,

20 said alkyl, alkenyl, alkynyl, cycloalkyl, aryl, and heterocyclyl is optionally substituted with up to three substituents selected from R^b , OH, (C_1-C_6) alkoxy, halogen, CO_2H , CN, $O(C=O)C_1-C_6$ alkyl, oxo, and $N(R^b)_2$;

25 R^a is (C_1-C_6) alkyl, (C_2-C_6) alkenyl, (C_2-C_6) alkynyl, (C_3-C_6) cycloalkyl, substituted or unsubstituted aryl, (C_1-C_6) perfluoroalkyl, 2,2,2-trifluoroethyl, or substituted or unsubstituted heterocyclyl; and

R^b is H, (C_1-C_6) alkyl, aryl, heterocyclyl, (C_3-C_6) cycloalkyl, $(C=O)OC_1-C_6$ alkyl, $(C=O)C_1-C_6$ alkyl or $S(O)_2R^a$;

30

R^c is selected from:

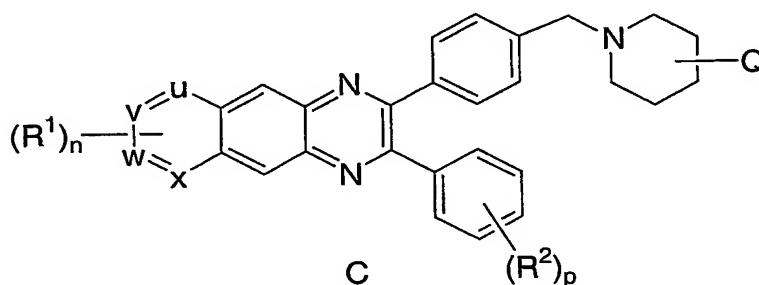
- 1) H,
- 2) C_1-C_{10} alkyl,
- 3) aryl,

- 4) C₂-C₁₀ alkenyl,
- 5) C₂-C₁₀ alkynyl,
- 6) heterocyclyl,
- 7) C₃-C₈ cycloalkyl,
- 8) C₁-C₆ perfluoroalkyl,

said alkyl, cycloalkyl, aryl, heterocyclyl, alkenyl, and alkynyl is optionally substituted with one or more substituents selected from R^Z, or

or a pharmaceutically acceptable salt or a stereoisomer thereof.

3. The compound according to Claim 1 which is:



wherein:

- a is 0 or 1;
- b is 0 or 1;
- m is 0, 1 or 2;
- n is 0, 1 or 2;
- p is 0, 1 or 2;
- r is 0 or 1;
- s is 0 or 1;
- u, v and x are independently selected from: CH and N;

w is selected from a bond, CH and N;

Q is selected from: -NR⁷R⁸ and heterocyclyl, said heterocyclyl optionally substituted with one to three substituents selected from R^Z;

R¹ is independently selected from:

- 1) (C=O)_aO_bC₁-C₁₀ alkyl,
- 2) (C=O)_aO_baryl,
- 3) C₂-C₁₀ alkenyl,
- 5 4) C₂-C₁₀ alkynyl,
- 5) (C=O)_aO_b heterocyclyl,
- 6) (C=O)_aO_bC₃-C₈ cycloalkyl,
- 7) CO₂H,
- 8) halo,
- 10 9) CN,
- 10) OH,
- 11) O_bC₁-C₆ perfluoroalkyl,
- 12) O_a(C=O)_bNR⁷R⁸,
- 13) NR^c(C=O)NR⁷R⁸,
- 15 14) S(O)_mR^a,
- 15) S(O)₂NR⁷R⁸,
- 16) NR^cS(O)_mR^a,
- 17) oxo,
- 18) CHO,
- 20 19) NO₂,
- 20) NR^c(C=O)O_bR^a,
- 21) O(C=O)O_bC₁-C₁₀ alkyl,
- 22) O(C=O)O_bC₃-C₈ cycloalkyl,
- 23) O(C=O)O_baryl, and
- 25 24) O(C=O)O_b-heterocycle,

said alkyl, aryl, alkenyl, alkynyl, heterocyclyl, and cycloalkyl optionally substituted with one or more substituents selected from R^z;

R² is independently selected from:

- 30 1) (C=O)_aO_bC₁-C₁₀ alkyl,
- 2) (C=O)_aO_baryl,
- 3) C₂-C₁₀ alkenyl,
- 4) C₂-C₁₀ alkynyl,
- 5) (C=O)_aO_b heterocyclyl,

- 6) $(\text{C}=\text{O})_a\text{O}_b\text{C}_3\text{-C}_8$ cycloalkyl,
 7) CO_2H ,
 8) halo,
 9) CN ,
 5 10) OH ,
 11) $\text{O}_b\text{C}_1\text{-C}_6$ perfluoroalkyl,
 12) $\text{O}_a(\text{C}=\text{O})_b\text{NR}^7\text{R}^8$,
 13) $\text{NR}^c(\text{C}=\text{O})\text{NR}^7\text{R}^8$,
 14) $\text{S}(\text{O})_m\text{R}^a$,
 10 15) $\text{S}(\text{O})_2\text{NR}^7\text{R}^8$,
 16) $\text{NR}^c\text{S}(\text{O})_m\text{R}^a$,
 17) CHO ,
 18) NO_2 ,
 19) $\text{NR}^c(\text{C}=\text{O})\text{O}_b\text{R}^a$,
 15 20) $\text{O}(\text{C}=\text{O})\text{O}_b\text{C}_1\text{-C}_{10}$ alkyl,
 21) $\text{O}(\text{C}=\text{O})\text{O}_b\text{C}_3\text{-C}_8$ cycloalkyl,
 22) $\text{O}(\text{C}=\text{O})\text{O}_b\text{aryl}$, and
 23) $\text{O}(\text{C}=\text{O})\text{O}_b\text{-heterocycle}$,
 said alkyl, aryl, alkenyl, alkynyl, heterocyclyl, and cycloalkyl optionally substituted
 20 with one, two or three substituents selected from R^z ;

R^7 and R^8 are independently selected from:

- 1) H ,
 2) $(\text{C}=\text{O})\text{O}_b\text{C}_1\text{-C}_{10}$ alkyl,
 25 3) $(\text{C}=\text{O})\text{O}_b\text{C}_3\text{-C}_8$ cycloalkyl,
 4) $(\text{C}=\text{O})\text{O}_b\text{aryl}$,
 5) $(\text{C}=\text{O})\text{O}_b\text{heterocyclyl}$,
 6) $\text{C}_1\text{-C}_{10}$ alkyl,
 7) aryl,
 30 8) $\text{C}_2\text{-C}_{10}$ alkenyl,
 9) $\text{C}_2\text{-C}_{10}$ alkynyl,
 10) heterocyclyl,
 11) $\text{C}_3\text{-C}_8$ cycloalkyl,
 12) SO_2R^a , and

13) $(C=O)NR^b_2$,

said alkyl, cycloalkyl, aryl, heterocyclyl, alkenyl, and alkynyl is optionally substituted with one or more substituents selected from R^Z , or

- 5 R^7 and R^8 can be taken together with the nitrogen to which they are attached to form a monocyclic or bicyclic heterocycle with 5-7 members in each ring and optionally containing, in addition to the nitrogen, one or two additional heteroatoms selected from N, O and S, said monocyclic or bicyclic heterocycle optionally substituted with one or more substituents selected from R^Z ;

10

R^Z is selected from:

- 1) $(C=O)_rO_s(C_1-C_{10})alkyl$,
- 2) $O_r(C_1-C_3)perfluoroalkyl$,
- 3) $(C_0-C_6)alkylene-S(O)_mR^a$,
- 15 4) oxo,
- 5) OH,
- 6) halo,
- 7) CN,
- 8) $(C=O)_rO_s(C_2-C_{10})alkenyl$,
- 20 9) $(C=O)_rO_s(C_2-C_{10})alkynyl$,
- 10) $(C=O)_rO_s(C_3-C_6)cycloalkyl$,
- 11) $(C=O)_rO_s(C_0-C_6)alkylene-aryl$,
- 12) $(C=O)_rO_s(C_0-C_6)alkylene-heterocyclyl$,
- 13) $(C=O)_rO_s(C_0-C_6)alkylene-N(R^b)_2$,
- 25 14) $C(O)R^a$,
- 15) $(C_0-C_6)alkylene-CO_2R^a$,
- 16) $C(O)H$,
- 17) $(C_0-C_6)alkylene-CO_2H$,
- 18) $C(O)N(R^b)_2$,
- 30 19) $S(O)_mR^a$, and
- 20) $S(O)_2NR^9R^{10}$
- 21) $NR^c(C=O)O_bR^a$,
- 22) $O(C=O)O_bC_1-C_{10}alkyl$,
- 23) $O(C=O)O_bC_3-C_8cycloalkyl$,

24) $\text{O}(\text{C}=\text{O})\text{O}_b\text{aryl}$, and

25) $\text{O}(\text{C}=\text{O})\text{O}_b\text{-heterocycle}$,

said alkyl, alkenyl, alkynyl, cycloalkyl, aryl, and heterocyclyl is optionally substituted with up to three substituents selected from R^b , OH, $(\text{C}_1\text{-C}_6)\text{alkoxy}$, halogen, CO_2H ,

5 CN , $\text{O}(\text{C}=\text{O})\text{C}_1\text{-C}_6$ alkyl, oxo, and $\text{N}(\text{R}^b)_2$;

R^a is $(\text{C}_1\text{-C}_6)\text{alkyl}$, $(\text{C}_2\text{-C}_6)\text{alkenyl}$, $(\text{C}_2\text{-C}_6)\text{alkynyl}$, $(\text{C}_3\text{-C}_6)\text{cycloalkyl}$, substituted or unsubstituted aryl, $(\text{C}_1\text{-C}_6)\text{perfluoroalkyl}$, 2,2,2-trifluoroethyl, or substituted or unsubstituted heterocyclyl; and

10

R^b is H, $(\text{C}_1\text{-C}_6)\text{alkyl}$, aryl, heterocyclyl, $(\text{C}_3\text{-C}_6)\text{cycloalkyl}$, $(\text{C}=\text{O})\text{OC}_1\text{-C}_6$ alkyl, $(\text{C}=\text{O})\text{C}_1\text{-C}_6$ alkyl or $\text{S}(\text{O})_2\text{R}^a$;

R^c is selected from:

15

1) H,

2) $\text{C}_1\text{-C}_{10}$ alkyl,

3) aryl,

4) $\text{C}_2\text{-C}_{10}$ alkenyl,

5) $\text{C}_2\text{-C}_{10}$ alkynyl,

20

6) heterocyclyl,

7) $\text{C}_3\text{-C}_8$ cycloalkyl,

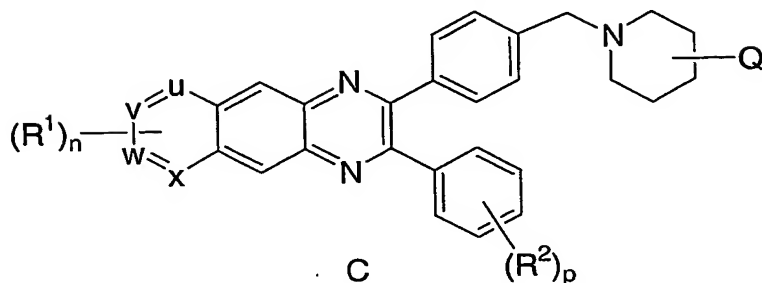
8) $\text{C}_1\text{-C}_6$ perfluoroalkyl,

said alkyl, cycloalkyl, aryl, heterocyclyl, alkenyl, and alkynyl is optionally substituted with one or more substituents selected from R^z , or

25

or a pharmaceutically acceptable salt or a stereoisomer thereof.

4. The compound according to Claim 2 which is:



wherein:

5

a is 0 or 1;

b is 0 or 1;

m is 0, 1 or 2;

n is 0, 1 or 2;

10 p is 0, 1 or 2;

r is 0 or 1;

s is 0 or 1;

u, v and x are independently selected from: CH and N;

15

w is selected from a bond, CH and N;

Q is selected from: -NR⁷R⁸, phenyl, benzimidazolyl, benzimidazolonyl, quinolinyl and isoquinolinyl, said benzimidazolyl, benzimidazolonyl, quinolinyl and

20 isoquinolinyl optionally substituted with Rᶻ;

R¹ is independently selected from:

1) (C=O)ₐOᵇC₁-C₁₀ alkyl,

2) (C=O)ₐOᵇaryl,

25 3) C₂-C₁₀ alkenyl,

4) C₂-C₁₀ alkynyl,

5) (C=O)ₐOᵇ heterocyclyl,

6) (C=O)ₐOᵇC₃-C₈ cycloalkyl,

- 7) CO_2H ,
 8) halo,
 9) CN ,
 10) OH ,
 5 11) $\text{O}_b\text{C}_1\text{-C}_6$ perfluoroalkyl,
 12) $\text{O}_a(\text{C}=\text{O})_b\text{NR}^7\text{R}^8$,
 13) $\text{NR}^c(\text{C}=\text{O})\text{NR}^7\text{R}^8$,
 14) $\text{S}(\text{O})_m\text{R}^a$,
 15) $\text{S}(\text{O})_2\text{NR}^7\text{R}^8$,
 10 16) $\text{NR}^c\text{S}(\text{O})_m\text{R}^a$,
 17) oxo,
 18) CHO ,
 19) NO_2 ,
 20) $\text{NR}^c(\text{C}=\text{O})\text{O}_b\text{R}^a$,
 15 21) $\text{O}(\text{C}=\text{O})\text{O}_b\text{C}_1\text{-C}_{10}$ alkyl,
 22) $\text{O}(\text{C}=\text{O})\text{O}_b\text{C}_3\text{-C}_8$ cycloalkyl,
 23) $\text{O}(\text{C}=\text{O})\text{O}_b$ aryl, and
 24) $\text{O}(\text{C}=\text{O})\text{O}_b$ -heterocycle,

said alkyl, aryl, alkenyl, alkynyl, heterocyclyl, and cycloalkyl optionally substituted
 20 with one or more substituents selected from R^Z ;

R^2 is independently selected from:

- 1) $(\text{C}=\text{O})_a\text{O}_b\text{C}_1\text{-C}_{10}$ alkyl,
 2) $(\text{C}=\text{O})_a\text{O}_b$ aryl,
 25 3) $\text{C}_2\text{-C}_{10}$ alkenyl,
 4) $\text{C}_2\text{-C}_{10}$ alkynyl,
 5) $(\text{C}=\text{O})_a\text{O}_b$ heterocyclyl,
 6) $(\text{C}=\text{O})_a\text{O}_b\text{C}_3\text{-C}_8$ cycloalkyl,
 7) CO_2H ,
 30 8) halo,
 9) CN ,
 10) OH ,
 11) $\text{O}_b\text{C}_1\text{-C}_6$ perfluoroalkyl,
 12) $\text{O}_a(\text{C}=\text{O})_b\text{NR}^7\text{R}^8$,

- 13) $\text{NR}^c(\text{C}=\text{O})\text{NR}^7\text{R}^8$,
 14) $\text{S}(\text{O})_m\text{R}^a$,
 15) $\text{S}(\text{O})_2\text{NR}^7\text{R}^8$,
 16) $\text{NR}^c\text{S}(\text{O})_m\text{R}^a$,
 5 17) CHO ,
 18) NO_2 ,
 19) $\text{NR}^c(\text{C}=\text{O})\text{O}_b\text{R}^a$,
 20) $\text{O}(\text{C}=\text{O})\text{O}_b\text{C}_1\text{-C}_{10}$ alkyl,
 21) $\text{O}(\text{C}=\text{O})\text{O}_b\text{C}_3\text{-C}_8$ cycloalkyl,
 10 22) $\text{O}(\text{C}=\text{O})\text{O}_b\text{aryl}$, and
 23) $\text{O}(\text{C}=\text{O})\text{O}_b\text{-heterocycle}$,

said alkyl, aryl, alkenyl, alkynyl, heterocyclyl, and cycloalkyl optionally substituted with one, two or three substituents selected from R^z ;

15 R^7 and R^8 are independently selected from:

- 1) H,
 2) $(\text{C}=\text{O})\text{O}_b\text{C}_1\text{-C}_{10}$ alkyl,
 3) $(\text{C}=\text{O})\text{O}_b\text{C}_3\text{-C}_8$ cycloalkyl,
 4) $(\text{C}=\text{O})\text{O}_b\text{aryl}$,
 20 5) $(\text{C}=\text{O})\text{O}_b\text{heterocyclyl}$,
 6) $\text{C}_1\text{-C}_{10}$ alkyl,
 7) aryl,
 8) $\text{C}_2\text{-C}_{10}$ alkenyl,
 9) $\text{C}_2\text{-C}_{10}$ alkynyl,
 25 10) heterocyclyl,
 11) $\text{C}_3\text{-C}_8$ cycloalkyl,
 12) SO_2R^a , and
 13) $(\text{C}=\text{O})\text{NR}^b_2$,

30 said alkyl, cycloalkyl, aryl, heterocyclyl, alkenyl, and alkynyl is optionally substituted with one or more substituents selected from R^z , or

R^7 and R^8 can be taken together with the nitrogen to which they are attached to form a monocyclic or bicyclic heterocycle with 5-7 members in each ring and optionally containing, in addition to the nitrogen, one or two additional heteroatoms selected

from N, O and S, said monocyclic or bicyclic heterocycle optionally substituted with one or more substituents selected from R^Z;

R^Z is selected from:

- 1) (C=O)_rO_s(C₁-C₁₀)alkyl,
- 5 2) O_r(C₁-C₃)perfluoroalkyl,
- 3) (C₀-C₆)alkylene-S(O)_mR^a,
- 4) oxo,
- 5) OH,
- 6) halo,
- 10 7) CN,
- 8) (C=O)_rO_s(C₂-C₁₀)alkenyl,
- 9) (C=O)_rO_s(C₂-C₁₀)alkynyl,
- 10) (C=O)_rO_s(C₃-C₆)cycloalkyl,
- 11) (C=O)_rO_s(C₀-C₆)alkylene-aryl,
- 15 12) (C=O)_rO_s(C₀-C₆)alkylene-heterocyclyl,
- 13) (C=O)_rO_s(C₀-C₆)alkylene-N(R^b)₂,
- 14) C(O)R^a,
- 15) (C₀-C₆)alkylene-CO₂R^a,
- 16) C(O)H,
- 20 17) (C₀-C₆)alkylene-CO₂H,
- 18) C(O)N(R^b)₂,
- 19) S(O)_mR^a,
- 20) S(O)₂NR⁹R¹⁰
- 21) NRC(C=O)O_bR^a,
- 25 22) O(C=O)O_bC₁-C₁₀ alkyl,
- 23) O(C=O)O_bC₃-C₈ cycloalkyl,
- 24) O(C=O)O_baryl, and
- 25) O(C=O)O_b-heterocycle,

30 said alkyl, alkenyl, alkynyl, cycloalkyl, aryl, and heterocyclyl is optionally substituted with up to three substituents selected from R^b, OH, (C₁-C₆)alkoxy, halogen, CO₂H, CN, O(C=O)C₁-C₆ alkyl, oxo, and N(R^b)₂;

R^a is (C₁-C₆)alkyl, (C₃-C₆)cycloalkyl, aryl, or heterocyclyl; and

R^b is H, (C₁-C₆)alkyl, aryl, heterocyclyl, (C₃-C₆)cycloalkyl, (C=O)OC₁-C₆ alkyl, (C=O)C₁-C₆ alkyl or S(O)₂R^a;

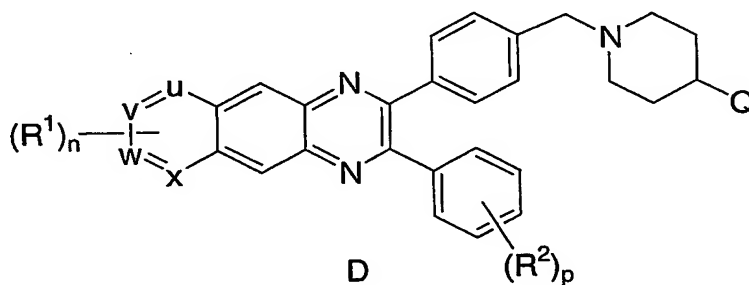
R^c is selected from:

- 1) H,
- 5 2) C₁-C₁₀ alkyl,
- 3) aryl,
- 4) C₂-C₁₀ alkenyl,
- 5) C₂-C₁₀ alkynyl,
- 6) heterocyclyl,
- 10 7) C₃-C₈ cycloalkyl,
- 8) C₁-C₆ perfluoroalkyl,

said alkyl, cycloalkyl, aryl, heterocyclyl, alkenyl, and alkynyl is optionally substituted with one or more substituents selected from R^z, or

15 or a pharmaceutically acceptable salt or a stereoisomer thereof.

5. The compound according to Claim 4 of the Formula D:



wherein

20 a is 0 or 1;

b is 0 or 1;

m is 0, 1 or 2;

n is 0, 1 or 2;

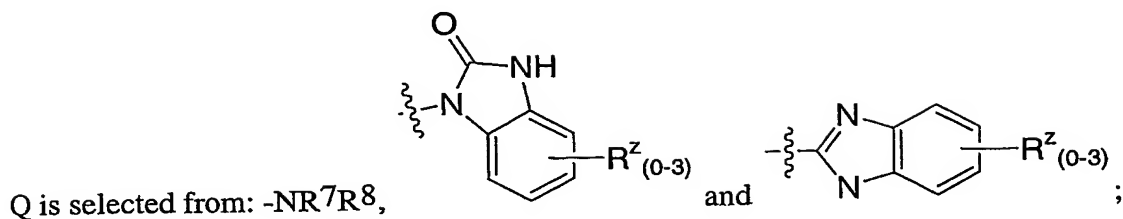
p is 0, 1 or 2;

25 r is 0 or 1;

s is 0 or 1;

u, v and x are independently selected from: CH and N;

w is selected from a bond, CH and N;



R¹ is independently selected from:

- 1) (C=O)_aO_bC₁-C₁₀ alkyl,
- 5 2) (C=O)_aO_baryl,
- 3) C₂-C₁₀ alkenyl,
- 4) C₂-C₁₀ alkynyl,
- 5) (C=O)_aO_b heterocyclyl,
- 6) (C=O)_aO_bC₃-C₈ cycloalkyl,
- 10 7) CO₂H,
- 8) halo,
- 9) CN,
- 10) OH,
- 11) O_bC₁-C₆ perfluoroalkyl,
- 15 12) O_a(C=O)_bNR⁷R⁸,
- 13) NR^c(C=O)NR⁷R⁸,
- 14) S(O)_mR^a,
- 15) S(O)₂NR⁷R⁸,
- 16) NR^cS(O)_mR^a,
- 20 17) oxo,
- 18) CHO,
- 19) NO₂,
- 20) NR^c(C=O)O_bR^a,
- 21) O(C=O)O_bC₁-C₁₀ alkyl,
- 25 22) O(C=O)O_bC₃-C₈ cycloalkyl,
- 23) O(C=O)O_baryl, and
- 24) O(C=O)O_b-heterocycle,

said alkyl, aryl, alkenyl, alkynyl, heterocyclyl, and cycloalkyl optionally substituted with one or more substituents selected from R^Z;

R² is independently selected from:

- 1) C₁-C₆ alkyl,
 - 2) aryl,
 - 3) heterocyclyl,
 - 5 4) CO₂H,
 - 5) halo,
 - 6) CN,
 - 7) OH,
 - 8) S(O)₂NR⁷R⁸,
- 10 said alkyl, aryl and heterocyclyl optionally substituted with one, two or three substituents selected from R^Z;

R⁷ and R⁸ are independently selected from:

- 1) H,
- 15 2) (C=O)O_bC₁-C₁₀ alkyl,
- 3) (C=O)O_bC₃-C₈ cycloalkyl,
- 4) (C=O)O_baryl,
- 5) (C=O)O_bheterocyclyl,
- 6) C₁-C₁₀ alkyl,
- 20 7) aryl,
- 8) C₂-C₁₀ alkenyl,
- 9) C₂-C₁₀ alkynyl,
- 10) heterocyclyl,
- 11) C₃-C₈ cycloalkyl,
- 25 12) SO₂R^a, and
- 13) (C=O)NR^b₂,

said alkyl, cycloalkyl, aryl, heterocyclyl, alkenyl, and alkynyl is optionally substituted with one or more substituents selected from R^Z, or

- 30 R⁷ and R⁸ can be taken together with the nitrogen to which they are attached to form a monocyclic or bicyclic heterocycle with 5-7 members in each ring and optionally containing, in addition to the nitrogen, one or two additional heteroatoms selected from N, O and S, said monocyclic or bicyclic heterocycle optionally substituted with one or more substituents selected from R^Z;

RZ is selected from:

- 1) $(\text{C}=\text{O})_r\text{O}_s(\text{C}_1\text{-C}_{10})\text{alkyl}$,
- 2) $\text{O}_r(\text{C}_1\text{-C}_3)\text{perfluoroalkyl}$,
- 3) $(\text{C}_0\text{-C}_6)\text{alkylene-S}(\text{O})_m\text{R}^a$,
- 5 4) oxo,
- 5) OH,
- 6) halo,
- 7) CN,
- 8) $(\text{C}=\text{O})_r\text{O}_s(\text{C}_2\text{-C}_{10})\text{alkenyl}$,
- 10 9) $(\text{C}=\text{O})_r\text{O}_s(\text{C}_2\text{-C}_{10})\text{alkynyl}$,
- 10) $(\text{C}=\text{O})_r\text{O}_s(\text{C}_3\text{-C}_6)\text{cycloalkyl}$,
- 11) $(\text{C}=\text{O})_r\text{O}_s(\text{C}_0\text{-C}_6)\text{alkylene-aryl}$,
- 12) $(\text{C}=\text{O})_r\text{O}_s(\text{C}_0\text{-C}_6)\text{alkylene-heterocyclyl}$,
- 13) $(\text{C}=\text{O})_r\text{O}_s(\text{C}_0\text{-C}_6)\text{alkylene-N}(\text{R}^b)_2$,
- 15 14) $\text{C}(\text{O})\text{R}^a$,
- 15) $(\text{C}_0\text{-C}_6)\text{alkylene-CO}_2\text{R}^a$,
- 16) $\text{C}(\text{O})\text{H}$,
- 17) $(\text{C}_0\text{-C}_6)\text{alkylene-CO}_2\text{H}$,
- 18) $\text{C}(\text{O})\text{N}(\text{R}^b)_2$,
- 20 19) $\text{S}(\text{O})_m\text{R}^a$,
- 20) $\text{S}(\text{O})_2\text{N}(\text{R}^b)_2$,
- 21) $\text{NR}^c(\text{C}=\text{O})\text{O}_b\text{R}^a$,
- 22) $\text{O}(\text{C}=\text{O})\text{O}_b\text{C}_1\text{-C}_{10}\text{ alkyl}$,
- 23) $\text{O}(\text{C}=\text{O})\text{O}_b\text{C}_3\text{-C}_8\text{ cycloalkyl}$,
- 25 24) $\text{O}(\text{C}=\text{O})\text{O}_b\text{aryl}$, and
- 25) $\text{O}(\text{C}=\text{O})\text{O}_b\text{-heterocycle}$,

said alkyl, alkenyl, alkynyl, cycloalkyl, aryl, and heterocyclyl is optionally substituted with up to three substituents selected from R^b , OH, $(\text{C}_1\text{-C}_6)\text{alkoxy}$, halogen, CO_2H , CN, $\text{O}(\text{C}=\text{O})\text{C}_1\text{-C}_6\text{ alkyl}$, oxo, and $\text{N}(\text{R}^b)_2$;

30

R^a is $(\text{C}_1\text{-C}_6)\text{alkyl}$, $(\text{C}_3\text{-C}_6)\text{cycloalkyl}$, aryl, or heterocyclyl; and

R^b is H, $(\text{C}_1\text{-C}_6)\text{alkyl}$, aryl, heterocyclyl, $(\text{C}_3\text{-C}_6)\text{cycloalkyl}$, $(\text{C}=\text{O})\text{OC}_1\text{-C}_6\text{ alkyl}$, $(\text{C}=\text{O})\text{C}_1\text{-C}_6\text{ alkyl}$ or $\text{S}(\text{O})_2\text{R}^a$;

R^C is selected from:

- 1) H,
- 2) C₁-C₁₀ alkyl,
- 5 3) aryl,
- 4) C₂-C₁₀ alkenyl,
- 5) C₂-C₁₀ alkynyl,
- 6) heterocyclyl,
- 7) C₃-C₈ cycloalkyl,
- 10 8) C₁-C₆ perfluoroalkyl,

said alkyl, cycloalkyl, aryl, heterocyclyl, alkenyl, and alkynyl is optionally substituted with one or more substituents selected from R^Z, or

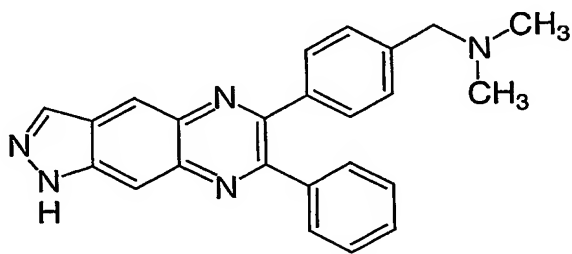
or a pharmaceutically acceptable salt or a stereoisomer thereof.

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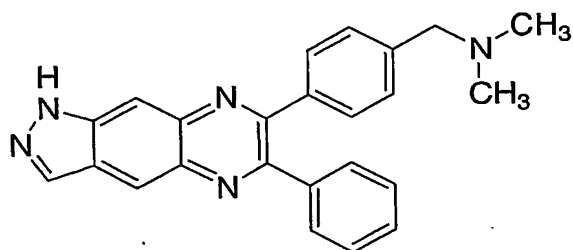
6. The TFA salt of a compound according to Claim 1 which is selected from:

- 1-{1-[4-(7-phenyl-1H-imidazo[4,5-g]quinoxalin-6-yl)benzyl]piperidin-4-yl}-1,3-dihydro-2H-benzimidazol-2-one;
- 20 N,N-dimethyl-1-[4-(6-phenyl-1H-imidazo[4,5-g]quinoxalin-7-yl)phenyl]metanamine;
- 1-{1-[4-(3-phenylbenzo[g]quinoxalin-2-yl)benzyl]piperidin-4-yl}-1,3-dihydro-2H-benzimidazol-2-one;
- 25 N-[(3R)-1-[4-(7-phenyl-1H-imidazo[4,5-g]quinoxalin-6-yl)benzyl]pyrrolidin-3-yl]-1,3-thiazole-5-carboxamide;
- 30 tert-butyl 1-[4-(7-phenyl-1H-imidazo[4,5-g]quinoxalin-6-yl)benzyl]azetidin-3-ylcarbamate;
- 9-{1-[4-(7-phenyl-1H-imidazo[4,5-g]quinoxalin-6-yl)benzyl]piperidin-4-yl}-9H-purin-6-amine;
- 35

6-(4-{[4-(3H-imidazo[4,5-b]pyridin-3-yl)piperidin-1-yl]methyl}phenyl)-7-phenyl-1H-imidazo[4,5-g]quinoxaline;

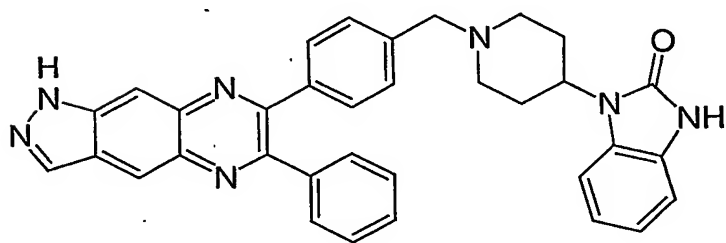


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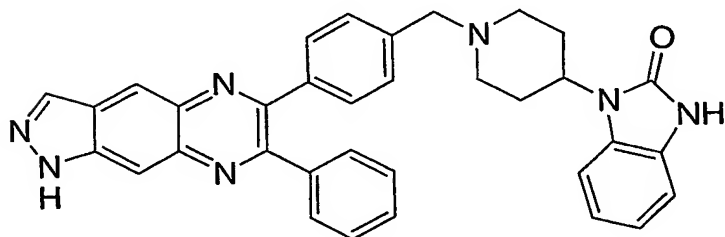


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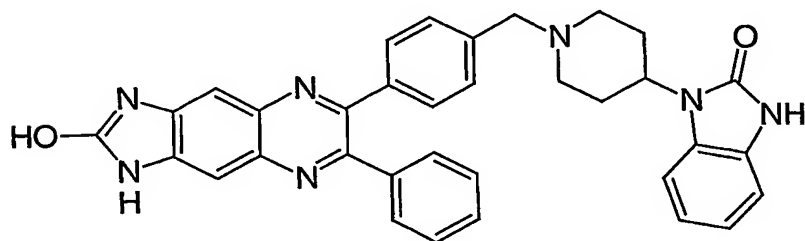
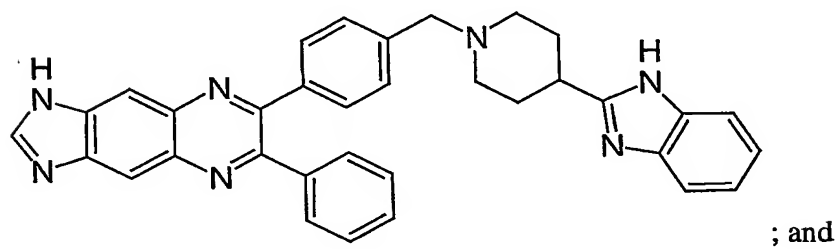
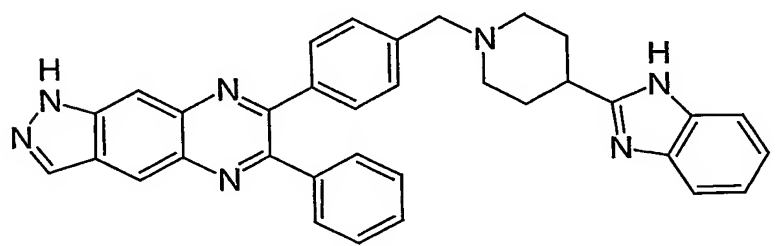
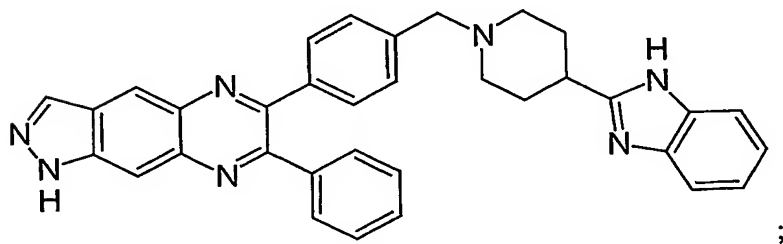
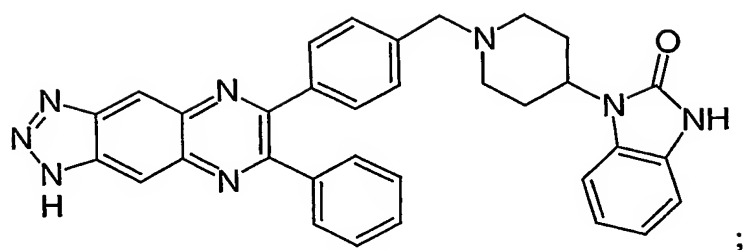
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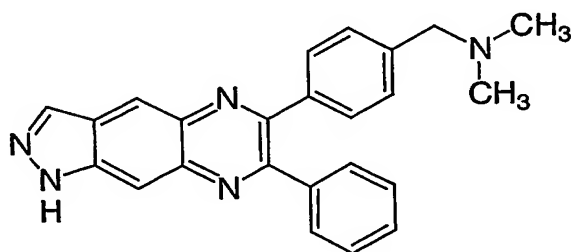


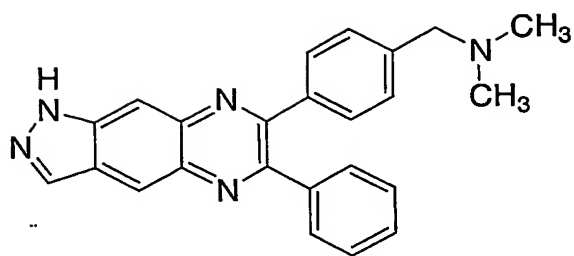
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or a stereoisomer thereof.

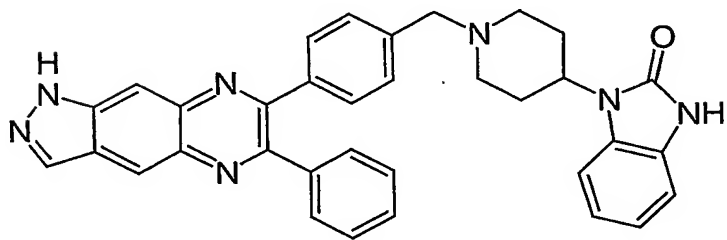
7. A compound which is selected from:

- 1-{1-[4-(7-phenyl-1H-imidazo[4,5-g]quinoxalin-6-yl)benzyl]piperidin-4-yl}-1,3-dihydro-2H-benzimidazol-2-one;
- 5 N,N-dimethyl-1-[4-(6-phenyl-1H-imidazo[4,5-g]quinoxalin-7-yl)phenyl]metanamine;
- 1-{1-[4-(3-phenylbenzo[g]quinoxalin-2-yl)benzyl]piperidin-4-yl}-1,3-dihydro-2H-benzimidazol-2-one;
- 10 N-[(3R)-1-[4-(7-phenyl-1H-imidazo[4,5-g]quinoxalin-6-yl)benzyl]pyrrolidin-3-yl]-1,3-thiazole-5-carboxamide;
- 15 tert-butyl 1-[4-(7-phenyl-1H-imidazo[4,5-g]quinoxalin-6-yl)benzyl]azetidin-3-ylcarbamate;
- 9-{1-[4-(7-phenyl-1H-imidazo[4,5-g]quinoxalin-6-yl)benzyl]piperidin-4-yl}-9H-purin-6-amine;
- 20 6-(4-{[4-(3H-imidazo[4,5-b]pyridin-3-yl)piperidin-1-yl]methyl}phenyl)-7-phenyl-1H-imidazo[4,5-g]quinoxaline;

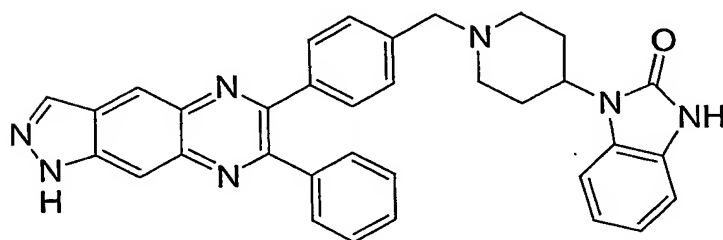




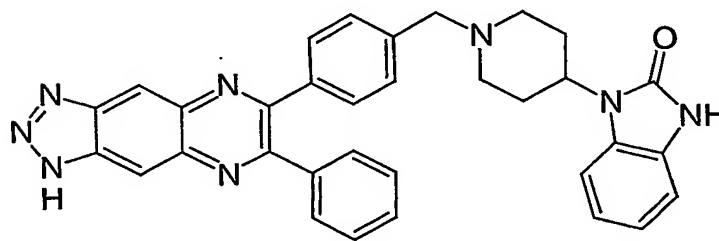
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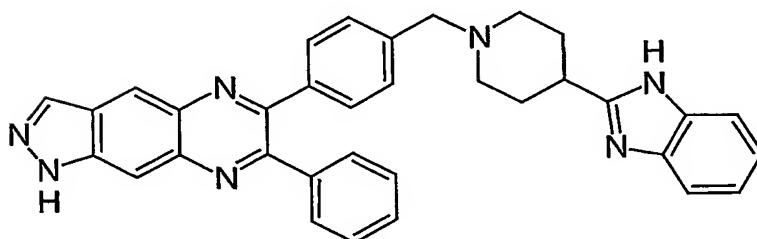
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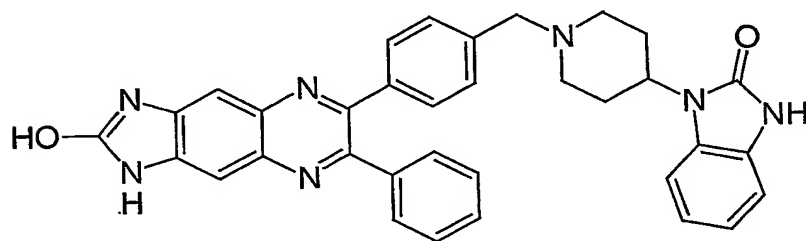
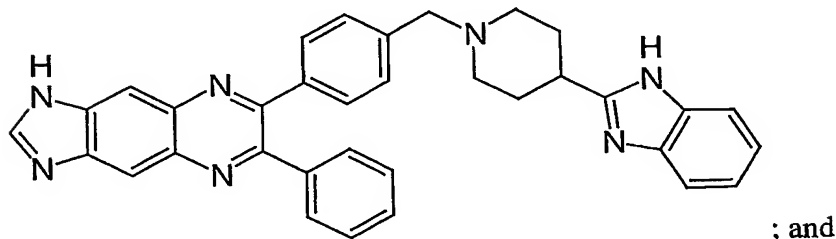
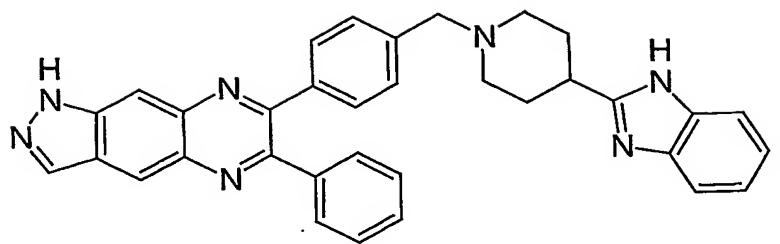
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or a pharmaceutically acceptable salt or a stereoisomer thereof.

5

8. A pharmaceutical composition comprising a pharmaceutical carrier, and dispersed therein, a therapeutically effective amount of a compound of Claim 1.

10

9. A pharmaceutical composition comprising a pharmaceutical carrier, and dispersed therein, a therapeutically effective amount of a compound of Claim 7.

15

10. A method of inhibiting one or more of the isoforms of Akt in a mammal which comprises administering to the mammal a therapeutically effective amount of a compound of Claim 1.

11. A method of inhibiting one or more of the isoforms of Akt in a mammal which comprises administering to the mammal a therapeutically effective amount of a compound of Claim 7.

5 12. A method for treating cancer which comprises administering to a mammal in need thereof a therapeutically effective amount of a compound of Claim 1.

10 13. A method for treating cancer which comprises administering to a mammal in need thereof a therapeutically effective amount of a compound of Claim 7.

15 14. A pharmaceutical composition made by combining the compound of Claim 1 and a pharmaceutically acceptable carrier.

15 15. A process for making a pharmaceutical composition comprising combining a compound of Claim 1 and a pharmaceutically acceptable carrier.

20 16. The composition of Claim 8 further comprising a second compound selected from:

- 1) an estrogen receptor modulator,
- 2) an androgen receptor modulator,
- 3) retinoid receptor modulator,
- 25 4) a cytotoxic agent,
- 5) an antiproliferative agent,
- 6) a prenyl-protein transferase inhibitor,
- 7) an HMG-CoA reductase inhibitor,
- 8) an HIV protease inhibitor,
- 30 9) a reverse transcriptase inhibitor,
- 10) an angiogenesis inhibitor,
- 11) a PPAR- γ agonists,
- 12) a PPAR- δ agonists,
- 13) an inhibitor of cell proliferation and survival signaling, and
- 35 14) an agent that interferes with a cell cycle checkpoint.

17. The composition of Claim 16, wherein the second compound is an angiogenesis inhibitor selected from the group consisting of a tyrosine kinase inhibitor, an inhibitor of epidermal-derived growth factor, an inhibitor of fibroblast-derived growth factor, an inhibitor of platelet derived growth factor, an MMP inhibitor, an integrin blocker, interferon- α , interleukin-12, pentosan polysulfate, a cyclooxygenase inhibitor, carboxyamidotriazole, combretastatin A-4, squalamine, 6-O-(chloroacetyl-carbonyl)-fumagillol, thalidomide, angiostatin and troponin-1.

18. The composition of Claim 16, wherein the second compound is an estrogen receptor modulator selected from tamoxifen and raloxifene.

19. A method of treating cancer which comprises administering a therapeutically effective amount of a compound of Claim 1 in combination with radiation therapy.

20. A method of treating or preventing cancer which comprises administering a therapeutically effective amount of a compound of Claim 1 in combination with a compound selected from:

- 1) an estrogen receptor modulator,
- 2) an androgen receptor modulator,
- 3) retinoid receptor modulator,
- 4) a cytotoxic agent,
- 5) an antiproliferative agent,
- 6) a prenyl-protein transferase inhibitor,
- 7) an HMG-CoA reductase inhibitor,
- 8) an HIV protease inhibitor,
- 9) a reverse transcriptase inhibitor,
- 10) an angiogenesis inhibitor,
- 11) a PPAR- γ agonists,
- 12) a PPAR- δ agonists,
- 13) an inhibitor of inherent multidrug resistance,
- 14) an anti-emetic agent,
- 15) an agent useful in the treatment of anemia,
- 16) an agent useful in the treatment of neutropenia,

- 17) an immunologic-enhancing drug,
- 18) an inhibitor of cell proliferation and survival signaling, and
- 19) an agent that interferes with a cell cycle checkpoint.

5 21. A method of treating cancer which comprises administering a therapeutically effective amount of a compound of Claim 1 in combination with radiation therapy and a compound selected from:

- 1) an estrogen receptor modulator,
- 2) an androgen receptor modulator,
- 10 3) retinoid receptor modulator,
- 4) a cytotoxic agent,
- 5) an antiproliferative agent,
- 6) a prenyl-protein transferase inhibitor,
- 7) an HMG-CoA reductase inhibitor,
- 15 8) an HIV protease inhibitor,
- 9) a reverse transcriptase inhibitor,
- 10) an angiogenesis inhibitor,
- 11) a PPAR- γ agonists,
- 12) a PPAR- δ agonists,
- 20 13) an inhibitor of inherent multidrug resistance,
- 14) an anti-emetic agent,
- 15) an agent useful in the treatment of anemia,
- 16) an agent useful in the treatment of neutropenia,
- 17) an immunologic-enhancing drug,
- 25 18) an inhibitor of cell proliferation and survival signaling, and
- 19) an agent that interferes with a cell cycle checkpoint.

22. A method of treating or preventing cancer which comprises administering a therapeutically effective amount of a compound of Claim 1 and paclitaxel or trastuzumab.

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